## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

## 1. (Currently amended) A compound of formula

$$R^3$$
 $R^1$ 
 $R^2$ 
 $R^2$ 
 $R^3$ 
 $R^2$ 

<u>or a stereoisomer</u> or <del>including the stereoisomers and the</del> pharmaceutically acceptable addition salts salt thereof,

wherein:

R<sup>1</sup> is NR<sup>4</sup>R<sup>5</sup> or OR<sup>5</sup>;

 $R^2$  is  $C_{1-6}$ alkyl;  $C_{1-6}$ alkyloxy or  $C_{1-6}$ alkylthio;

 $R^3 \quad \text{is} \quad \text{hydrogen,} \quad C_{1\text{-6}} \\ \text{alkyl}, \quad C_{1\text{-6}} \\ \text{alkylsulfonyl,} \quad C_{1\text{-6}} \\ \text{alkylthio;} \\$ 

 $R^4 \quad \text{is hydrogen,} \quad \underline{\text{or}} \quad C_{1\text{-6}} \\ \text{alkyl,} \quad \underline{\text{mono- or }} \\ \text{di}(C_{3\text{-6}} \\ \text{eyeloalkyl,}) \\ \text{methyl,} \quad C_{3\text{-6}} \\ \text{alkenyl,} \quad \text{hydroxy} \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{exercises} \\ \text{or } \\ C_{1\text{-6}} \\ \text{alkyl} \\ \text{exercises} \\ \text{exercises}$ 

 $R^5$  is  $C_{1-8}$ alkyl,  $mono-or-di(C_{3-6}$ alkyleyeloalkyl)methyl,  $Ar^{\dagger}CH_{25}$ ,  $C_{3-6}$ alkenyl,  $C_{1-6}$ alkyl,  $or-di(C_{1-6}$ alkyl, thienylmethyl, furanylmethyl,  $C_{1-6}$ alkyl,  $mono-or-di(C_{1-6}$ alkyl)amino $C_{1-6}$ alkyl,  $di(C_{1-6}$ alkyl)amino,  $C_{1-6}$ alkylearbonyl $C_{1-6}$ alkyl,  $C_{1-6}$ alkyl substituted with imidazolyl; or a radical of formula - $Alk-O-CO-Ar^{\dagger}$ ;

or R<sup>4</sup> and R<sup>5</sup> taken together with the nitrogen atom to which they are attached may form a pyrrolidinyl, piperidinyl, homopiperidinyl or morpholinyl group, optionally substituted with C<sub>1-6</sub>alkyl or C<sub>1-6</sub>alkyloxyC<sub>1-6</sub>alkyl; and

Ar is phenyl; phenyl substituted with 1, 2 or 3 substituents independently selected from halo, C<sub>1-6</sub>alkyl, trifluoromethyl, hydroxy, eyano, C<sub>1-6</sub>alkyloxy, benzyloxy, C<sub>1-6</sub>alkylthio, nitro, amino and mono- and di(C<sub>1-6</sub>alkyl)amino; pyridinyl; pyridinyl substituted with 1, 2 or 3 substituents independently selected from halo, C<sub>1-6</sub>alkyl, trifluoromethyl, hydroxy, eyano, C<sub>1-6</sub>alkyloxy, benzyloxy, C<sub>1-6</sub>alkylthio, nitro, amino, and mono- or di(C<sub>1-6</sub>alkyl)amino and piperidinyl; and wherein said substituted phenyl may optionally be further substituted with one or more halogens;

Ar<sup>1</sup> is phenyl; phenyl substituted with 1, 2 or 3 substituents each independently selected from halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkyloxy, di(C<sub>1-6</sub>alkyl)aminoC<sub>1-6</sub>alkyl, trifluoromethyl and C<sub>1-6</sub>alkyl substituted with morpholinyl; or pyridinyl; and

Alk is C<sub>1-6</sub>alkanediyl;

with the proviso that 5-methyl-3-phenyl-7-(phenylmethoxy)-pyrazolo[1,5-a]-pyrimidine and 2,5-dimethyl-7-(methylamino)-3-phenyl-pyrazolo[1,5-a]-pyrimidine are is not included.

## 2-13. (Cancelled)

- 14. (New) A compound according to claim 1, wherein  $R^3$  is hydrogen,  $C_{1-6}$ alkyl or  $C_{1-6}$ alkylthio.
- 15. (New) A compound according to claim 14, wherein  $R^3$  is hydrogen, methyl or  $CH_3S$ -.
- 16. (New) A compound according to claim 1, wherein R<sup>4</sup> is hydrogen or C<sub>2-4</sub>alkyl.

- 17. (New) A compound according to claim 16, wherein R<sup>4</sup> is hydrogen or n-propyl.
- 18. (New) A compound according to claim 1, wherein Ar is phenyl substituted with 1, 2 or 3 substituents independently selected from halo and  $C_{1-6}$ alkyloxy; or Ar is pyridinyl substituted with 1, 2 or 3 substituents independently selected from halo,  $C_{1-6}$ alkyl and  $di(C_{1-6}$ alkyl)amino.
- 19. (New) A compound according to claim 18, wherein Ar is phenyl substituted with 2 or 3 substituents independently selected from halo and methoxy; or Ar is pyridinyl substituted with 2 or 3 substituents independently selected from halo, methyl and dimethylamino.
  - 20. (New) A compound according to claim 1, wherein R<sup>2</sup> is methyl.
- 21. (New) A compound according to claim 1, wherein  $R^3$  is hydrogen,  $C_{1-6}$ alkyl or  $C_{1-6}$ alkylthio;  $R^4$  is hydrogen or  $C_{2-4}$ alkyl; and Ar is phenyl substituted with 1, 2 or 3 substituents independently selected from halo and  $C_{1-6}$ alkyloxy, or pyridinyl substituted with 1, 2 or 3 substituents independently selected from halo,  $C_{1-6}$ alkyl and  $di(C_{1-6}$ alkyl)amino.
- 22. (New) A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and a therapeutically effective amount of a compound according to claim 1.
- 23. (New) A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and a therapeutically effective amount of a compound according to claim 15.

- 24. (New) A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and a therapeutically effective amount of a compound according to claim 17.
- 25. (New) A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and a therapeutically effective amount of a compound according to claim 19.
- 26. (New) A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and a therapeutically effective amount of a compound according to claim 21.
- 27. (New) A method for treating an endocrine, psychiatric or neurologic disorder or illness in a warm-blooded animal comprising administering to said animal in need of treatment a therapeutically effective amount of a compound according to claim 1.
- 28. (New) A method for treating an endocrine, psychiatric or neurologic disorder or illness in a warm-blooded animal comprising administering to said animal in need of treatment a therapeutically effective amount of a compound according to claim 18.
- 29. (New) A method for treating an endocrine, psychiatric or neurologic disorder or illness in a warm-blooded animal comprising administering to an animal in need thereof a therapeutically effective amount of a compound according to claim 21.
- 30. (New) A method for treating a physiological condition or disorder in a warm-blooded animal arising from the hyper-secretion of corticotrophin-releasing

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factor comprising administering to an animal in need thereof a therapeutically effective amount of a compound according to claim 1.

- 31. (New) A method for treating a physiological condition or disorder in a warm-blooded animal arising from the hyper-secretion of corticotrophin-releasing factor comprising administering to an animal in need thereof a therapeutically effective amount of a compound according to claim 18.
- 32. (New) A method for treating a physiological condition or disorder in a warm-blooded animal arising from the hyper-secretion of corticotrophin-releasing factor comprising administering to an animal in need thereof a therapeutically effective amount of a compound according to claim 21.

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